DYNAREX ANTIFUNGAL- antifungal cream
Dynarex Corporation

1231, 1233

Active Ingredient

Active Ingredient	Purpose
Clotrimazole 1.0%	Antifungal

Purpose

- For treatment of most athlete's foot (tinea pedis), jock itch (tinea crusis), Ringworm (tinea corporis).
- For the treatment of superficial skin infections caused by Yeast (Candida Albicans).
- Relieves itching, scaling, cracking, burning, redness, soreness, irritation discomfort and chafing associated with jock itch.

Warnings

Do not use:

- Do not use on children under 2 years of age unless directed by a doctor.
- Avoid contact with eyes.
- For athletes foot and ringworm - if irritation occurs, or if there is no improvement within 4 weeks, discontinue use and consult a doctor.
- For jock itch - if irritation occurs, or if there is no improvement within two weeks, discontinue use and consult a doctor.
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Indications and Usage

- Clean the affected area and dry thoroughly. Apply a layer of cream over affected area twice daily (morning and night) or as directed by a doctor.
- Supervise children in the use of this product.
- For athlete's foot, pay special attention to spaces between toes: wear well fitting, ventilated shoes, and change shoes and socks at least once daily.

Dosage and Administration

- Foe athlete's foot and ringworm, use daily for 4 weeks.
- For jock itch, use daily for 2 weeks.If conditions persist longer, consult a doctor.
- This product is not effective on scalp or nails.

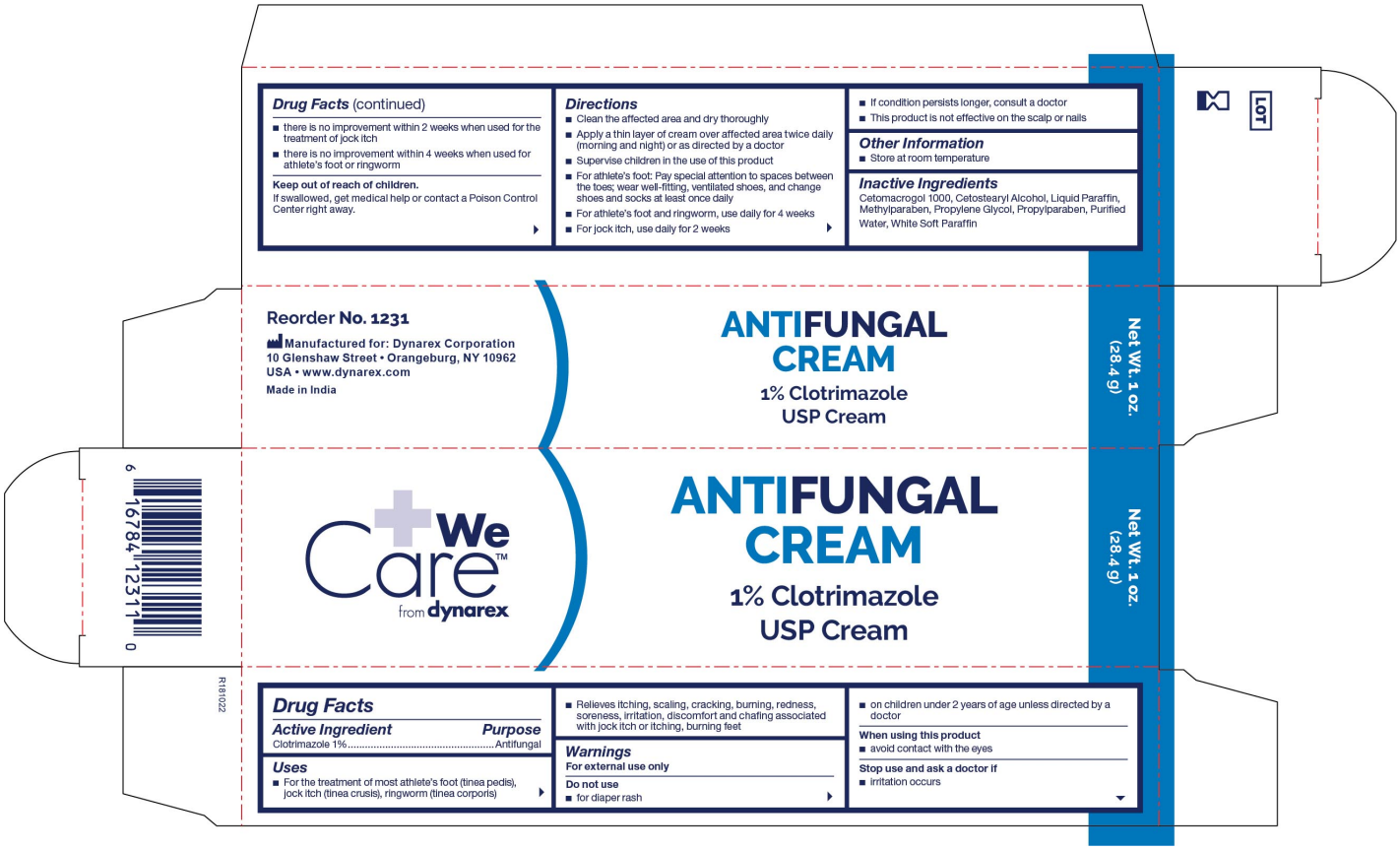
Keep Out Of Reach Of Children

KEEP OUT Of REACH Of CHILDREN

INACTIVE INGREDIENTS

Inactive Ingredients: Cetomicrogol 1000, Cetostearyl alcohol, Liquid paraffin, Methylparaben, Propylene glycol, Propylparaben, Purified water, White soft paraben

Label



Label

DYNAREX ANTIFUNGAL

antifungal cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-231
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
CETETH-20 (UNII: I835H2IHHX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-231-13	72 in 1 CASE	05/18/2010	
1	NDC:67777-231-12	1 in 1 BOX		
1	NDC:67777-231-02	28.25 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:67777-231-11	24 in 1 CASE	05/18/2010	
2	NDC:67777-231-10	6 in 1 BOX		
2	NDC:67777-231-01	113 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	05/18/2010	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 11/2024 Dynarex Corporation